## **AMENDMENTS**

## **Listing of Claims**

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- 1-29. (Canceled)
- 30. (Presently amended) A method <u>forof</u> screening <u>for</u> protein <u>stability</u>, folding and/or solubility mutants comprising:
  - a) providing a gene encoding <u>a</u> fusion protein comprising (i) a protein of interest and (ii) a first segment of a marker protein, wherein said first segment does not affect the folding or solubility of the protein of interest,—, wherein said fusion protein is <u>unstable</u>, not properly folded and/or <u>not</u> soluble when expressed in said host cell;
  - b) mutagenizing that portion of the gene encoding said protein of interest;
  - c) expressing said fusion protein in a host cell that expresses a second segment of said marker protein, wherein said second segment is capable of structural complementation with said first segment; and
  - d) determining structural complementation,

wherein a relative increase in structural complementation, as compared to the structural complementation observed with the unmutagenized fusion protein, indicates an increase in <u>stability</u>, proper folding and/or solubility of said protein.

- 31. (Original) The method of claim 30, wherein said fusion is C-terminal to said protein of interest.
- 32 (Original) The method of claim 30, wherein said fusion is N-terminal to said protein of interest.

- 33. (Original) The method of claim 30, wherein said marker protein is selected from the group consisting of a target binding protein, an enzyme, a protein inhibitor, a chromophore.
- 34. (Original) The method of claim 30, wherein said host cell is selected from the group consisting of a bacterial cell, an insect cell, a yeast cell, a nematode cell, a mammalian cell.

## 35-40. (Canceled)

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- 41. (New) The method of claim 30, wherein the unmutagenized fusion protein is unstable.
- 42. (New) The method of claim 30, wherein the unmutagenized fusion protein is not properly folded.
- 43. (New) The method of claim 30, wherein the unmutagenized fusion protein is not soluble.